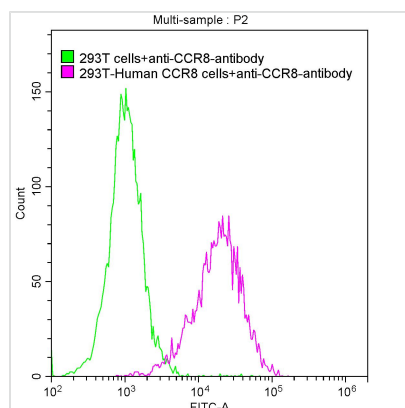




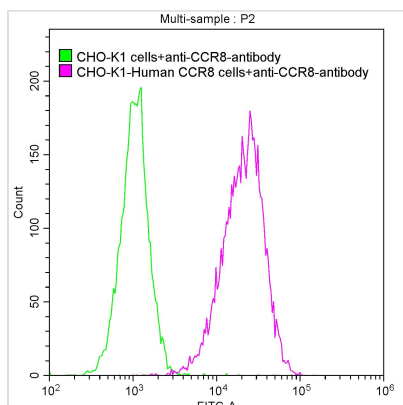
# CCR8 Recombinant Monoclonal Antibody

<b>Product Code</b>	CSB-RA004847MA4HU
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	P51685
<b>Immunogen</b>	Recombinant Human CCR8 protein
<b>Species Reactivity</b>	Human
<b>Tested Applications</b>	FC
<b>Form</b>	Liquid
<b>Conjugate</b>	Non-conjugated
<b>Storage Buffer</b>	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4
<b>Purification Method</b>	Affinity-chromatography
<b>Isotype</b>	hIgG1?lambda 2
<b>Clonality</b>	Monoclonal
<b>Product Type</b>	Recombinant Antibody
<b>Immunogen Species</b>	Homo sapiens (Human)
<b>Research Area</b>	Immunology
<b>Gene Names</b>	CCR8
<b>Clone No.</b>	10A9

## Image



Untransfected HEK293T cells surface(green line) and transfected Human CCR8 HEK293T stable cells surface (red line) were stained with anti-CCR8 recombinant antibody (2μg/1\*10<sup>6</sup>cells), washed and then followed by FITC-conjugated anti-Human IgG Fc antibody and analyzed with flow cytometry.



Untransfected CHO-K1 cells surface (green line) and transfected Human CCR8 CHO-K1 stable cells surface (red line) were stained with anti-CCR8 recombinant antibody ( $2\mu\text{g}/1 \times 10^6$  cells), washed and then followed by FITC-conjugated anti-Human IgG Fc antibody and analyzed with flow cytometry.

## Description

CUSABIO meticulously generated the CCR8 recombinant monoclonal antibody through a systematic procedure. Initially, B cells were extracted from the immunized animal's spleen, employing the recombinant human CCR8 protein as the immunogen during the immunization process. Subsequently, RNA isolation from the B cells was followed by cDNA synthesis through reverse transcription. By utilizing the cDNA as a template, the gene encoding the CCR8 antibody was extended using a degenerate primer and inserted into a recombinant vector. The recombinant vector was then introduced into host cells via transfection, facilitating the expression of the CCR8 recombinant monoclonal antibodies. These antibodies were harvested from the cell culture supernatant and subjected to purification using affinity chromatography. Rigorous validation testing FC was conducted to verify this antibody's reactivity with human CCR8 protein, ensuring its specificity and reliability.